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NEWS	10	JAN	26	Updated MeSH vocabulary, new structured abstracts, and other enhancements improve searching in STN reload of MEDLINE
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NEWS	12	FEB	23	PCTFULL file on STN completely reloaded
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NEWS	14	FEB	25	LPCI will be replaced by LDPCI
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NEWS	17	APR	28	The DWPI (files WPINDEX, WPIDS and WPIX) on STN have been enhanced with thesauri for the European Patent Classifications
NEWS	18	MAY	02	MEDLINE Improvements Provide Fast and Simple Access to DOI and Chemical Name Information
NEWS	19	MAY	12	European Patent Classification thesauri added to the INPADOC files, PCTFULL, GBFULL and FRFULL
NEWS	20	MAY	23	Enhanced performance of STN biosequence searches
NEWS	21	MAY	23	Free Trial of the Numeric Property Search Feature in PCTFULL on STN
NEWS	22	JUN	20	STN on the Web Enhanced with New Patent Family Assistant and Updated Structure Plug-In
NEWS	23	JUN	20	INPADOC databases enhanced with first page images
NEWS		JUN		PATDPA database updates to end in June 2011
NEWS		JUN		INPADOC: Delay of German patent coverage
NEWS		JUN		MARPAT Enhancements Save Time and Increase Usability
NEWS		JUL		STN adds Australian patent full-text database,
NEWS		AUG		AUPATFULL, including the new numeric search feature. CA Sections Added to ACS Publications Web Editions Platform

NEWS EXPRESS 17 DECEMBER 2010 CURRENT WINDOWS VERSION IS V8.4.2 .1, AND CURRENT DISCOVER FILE IS DATED 24 JANUARY 2011.

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ENTRY SESSION
0.23 0.23

FULL ESTIMATED COST

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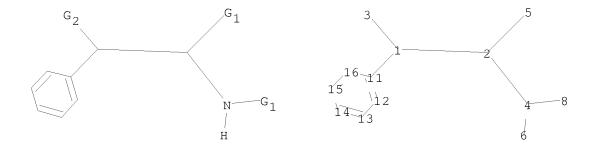
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chain nodes :
1 2 3 4 5 6 8
ring nodes :
11 12 13 14 15 16
chain bonds :
1-2 1-3 1-11 2-4 2-5 4-6 4-8
ring bonds :
11-12 11-16 12-13 13-14 14-15 15-16
exact/norm bonds :
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normalized bonds :
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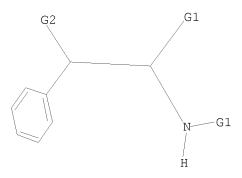
G1:Ak, COOH, H, Cb

G2:H,OH

Match level:
1:CLASS 2:CLASS 3:CLASS 4:CLASS 5:CLASS 6:CLASS 8:CLASS 11:Atom 12:Atom 13:Atom 14:Atom 15:Atom 16:Atom

L1 STRUCTURE UPLOADED

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G1:Ak,COOH,H,Cb G2:H,OH Structure attributes must be viewed using STN Express query preparation.

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FULL SEARCH INITIATED 18:24:31 FILE 'REGISTRY' FULL SCREEN SEARCH COMPLETED - 4925886 TO ITERATE

4.2% PROCESSED 206855 ITERATIONS 155190 ANSWERS

4.7% PROCESSED 231216 ITERATIONS

177661 ANSWERS

100.0% PROCESSED 4925886 ITERATIONS

2168476 ANSWERS

SEARCH TIME: 00.00.37

2168476 SEA SSS FUL L1 L2

=> file caplus

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197.37 197.60

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FILE COVERS 1907 - 4 Aug 2011 VOL 155 ISS 6
FILE LAST UPDATED: 3 Aug 2011 (20110803/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Jun 2011
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SINCE FILE TOTAL ENTRY SESSION 0.52 198.64

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=> s 12 and depression

4 DEPRESSION

L3 0 L2 AND DEPRESSION

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COST IN U.S. DOLLARS

SINCE FILE TOTAL ENTRY SESSION 6.72 205.36

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FILE COVERS 1907 - 4 Aug 2011 VOL 155 ISS 6
FILE LAST UPDATED: 3 Aug 2011 (20110803/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Jun 2011
USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Jun 2011

CAplus now includes complete International Patent Classification (IPC) reclassification data for the first quarter of 2011.

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This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s phenylalanine and depression

98709 PHENYLALANINE

106211 DEPRESSION

L4 459 PHENYLALANINE AND DEPRESSION

=> s 14 and carbonic anhydrase

61649 CARBONIC

15374 ANHYDRASE

15199 CARBONIC ANHYDRASE

(CARBONIC (W) ANHYDRASE)

L5 4 L4 AND CARBONIC ANHYDRASE

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L5 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2011 ACS on STN

ACCESSION NUMBER: 2008:9031 CAPLUS

DOCUMENT NUMBER: 148:113242

TITLE: Treatment of depressive disorders INVENTOR(S): Sun, Miao-Kun; Alkon, Daniel L.

PATENT ASSIGNEE(S): Blanchette Rockefeller Neurosciences Institute, USA

SOURCE: U.S. Pat. Appl. Publ., 20 pp., Cont.-in-part of U.S.

Ser. No. 594,420.

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

	PATENT NO.								APPLICATION NO.										
	US 20080003181				A1 2				US 2007-802724										
	US 7977377																		
	WO 2005115548								WO 2	005-1		20050518							
	WO	WO 2005115548			A3 20070322														
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ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT

OTHER SOURCE(S): MARPAT 148:113242

AB The invention provides for the use of carbonic anhydrase activators; protein kinase C activators and FGF-18 to treat depressive disorders. The invention also relates to improved animal models and methods for screening and identifying compds. the treatment of depressive disorders.

L5 ANSWER 2 OF 4 CAPLUS COPYRIGHT 2011 ACS on STN

ACCESSION NUMBER: 2005:1289949 CAPLUS

DOCUMENT NUMBER: 144:640

TITLE: Treatment of depressive disorders INVENTOR(S): Sun, Miao-Kun; Alkon, Daniel L.

PATENT ASSIGNEE(S): Blanchette Rockefeller Neurosciences Institute, USA

SOURCE: PCT Int. Appl., 40 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND DATE	E APPL:	ICATION NO.	DATE		
WO 2005115548	A2 2005	51208 WO 2	005-US17158	20050518		
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     US 20080003181
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     US 20090041667
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                                                                   20080630
PRIORITY APPLN. INFO.:
                                            US 2004-571892P
                                                                P 20040518
                                            EP 2005-749738
                                                                A3 20050518
                                            WO 2005-US17158
                                                                W 20050518
                                            US 2008-594420
                                                                A2 20080630
ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT
OTHER SOURCE(S):
                        MARPAT 144:640
    The invention provides for the use of carbonic anhydrase activators,
     protein kinase C activators and FGF-18 to treat depressive disorders.
     invention also relates to improved animal models and methods for screening
     and identifying compds. for the treatment of depressive disorders.
OS.CITING REF COUNT:
                               THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD
                         2
                               (2 CITINGS)
REFERENCE COUNT:
                         1
                               THERE ARE 1 CITED REFERENCES AVAILABLE FOR THIS
                               RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT
     ANSWER 3 OF 4 CAPLUS COPYRIGHT 2011 ACS on STN
ACCESSION NUMBER:
                         2003:535074 CAPLUS
DOCUMENT NUMBER:
                         139:270901
TITLE:
                         Carbonic anhydrase activators. The selective
                         serotonin reuptake inhibitors fluoxetine, sertraline
                         and citalopram are strong activators of isozymes I and
                         TT
                         Casini, Angela; Caccia, Silvio; Scozzafava, Andrea;
AUTHOR(S):
                         Supuran, Claudiu T.
                         Dipartimento di Chimica, Laboratorio di Chimica
CORPORATE SOURCE:
                         Bioinorganica, Universita degli Studi di Firenze,
                         Sesto Fiorentino (Firenze), I-50019, Italy
SOURCE:
                         Bioorganic & Medicinal
Chemistry Letters (2003),
                         13(16), 2765-2768
                         CODEN: BMCLE8; ISSN: 0960-894X
                         Elsevier Science B.V.
PUBLISHER:
DOCUMENT TYPE:
                         Journal
LANGUAGE:
                         English
     The selective serotonin reuptake inhibitors (SSRI) fluoxetine, sertraline
     and citalopram have been investigated for their ability to activate two
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carbonic anhydrase (CA) isoenzymes, hCA I and hCA II, in parallel with two standard activators for which the x-ray structure (in complex with

isoenzyme II) has been resolved: histamine and phenylalanine. All three SSRI activated both isoenzymes with potencies comparable to that of the stds. although the profile was different: for hCA I, best activators were fluoxetine and histamine, with citalopram and sertraline showing weaker activity. For hCA II, the best activators were phenylalanine and citalopram, and the weakest histamine and sertraline, whereas fluoxetine showed an intermediate behavior. These results suggest that SSRI efficacy in major depression complicating Alzheimer's disease may be partly due to their ability to activate CA isoenzymes and may lead to the development of potent activators for the therapy of diseases associated with significant decreases in brain CA activity.

OS.CITING REF COUNT: 8 THERE ARE 8 CAPLUS RECORDS THAT CITE THIS RECORD

(8 CITINGS)

REFERENCE COUNT: 35 THERE ARE 35 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 4 OF 4 CAPLUS COPYRIGHT 2011 ACS on STN

ACCESSION NUMBER: 2002:594810 CAPLUS

DOCUMENT NUMBER: 137:155177

TITLE: Preparation and ophthalmic compositions of amino acid

amides for treating ocular hypertension

INVENTOR(S): Garcia, Maria L.; Kaczorowski, Gregory J.; Gao,

Ying-Duo

PATENT ASSIGNEE(S): Merck & Co., Inc., USA SOURCE: PCT Int. Appl., 38 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

F	PAT	ENT 1	NO.			KIND DA			ATE APPLICATION AP			ION 1	N NO.			DATE			
	10	O 2002060863				A1		20020808		WO 2002-US3049					20020124				
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P	U	2002	2518	56		A1	L 20020812 AU 2002-251856							20020124					
J	JP 2004530647						T 20041007 JP 2002-561014						20020124						
E	EP 1358153					A1	20031105 EP 200)2-720887 2				20020125		
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ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT OTHER SOURCE(S): MARPAT 137:155177

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The compds. with a general formula of I [wherein R and R2 = independently AB alkyl, (CH2)n(hetero)aryl, (CH2)nheterocycloalkyl, said alkyl or (hetero)aryl optionally substituted with 1-3 groups of R3; Y =(CH2)nSCOR4; X = CH2 or O in which m = 1; R3 = H, alkoxy, alkyl(amino), CF3, NO2, NH2, CN, or halo; R4 = alkoxy or alkyl; R7 = H, halo, OH, NO2, NH2, CN, alkoxycarbonyl, CO2H, haloalkyl, alkoxycarbonylalkyl, or alkylsulfonyl; m = 1-3; n = 0-3; or a pharmaceutically acceptable salt, enantiomer, diastereomer, or mixture thereof] were prepared For example, L-leucine derivative II was prepared in a 7-step synthesis involving condensation of 4-phenyl-2-(acetylthiomethyl)butyric acid and (S)-leucine t-Bu ester and amidation with aniline (50%). This invention relates to the use of potent potassium channel blockers or formulations thereof in the treatment of glaucoma and other conditions which leads to elevated intraoccular pressure in the eye of a patient. This invention also relates to the use of such compds. to provide a neuroprotective effect to the eye of mammalian species, particularly humans. The compds. I were found to cause concentration dependent inhibition of the fluorescence ratio

with IC50 values in the range of 10 nM to 5 $\mu\text{M}\text{,}$ more preferably from 100 nM to 1 $\mu\text{M}\text{.}$

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(FILE 'HOME' ENTERED AT 18:23:58 ON 04 AUG 2011)

FILE 'REGISTRY' ENTERED AT 18:24:09 ON 04 AUG 2011 L1 STRUCTURE UPLOADED L2 2168476 S L1 FULL

FILE 'CAPLUS' ENTERED AT 18:25:16 ON 04 AUG 2011

FILE 'CAPLUS' ENTERED AT 18:25:57 ON 04 AUG 2011

FILE 'REGISTRY' ENTERED AT 18:26:12 ON 04 AUG 2011
0 S L2 AND DEPRESSION

FILE 'CAPLUS' ENTERED AT 18:27:37 ON 04 AUG 2011
459 S PHENYLALANINE AND DEPRESSION
L5 4 S L4 AND CARBONIC ANHYDRASE

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